



# EC Declaration of Conformity

We herewith declare, in exclusive responsibility, that the instrument

## **Leica ASP300 – Vacuum Tissue Processor**

was developed, designed and manufactured to conform with the

- Council Directive 73/23/EEC, (Low Voltage),
- Council Directive 89/336/EEC, Appendix I (Electromagnetic Compatibility) and
- European council Directive 98/79/EC (IVD)

including their amendments up to the date mentioned below.

The following harmonized standards were applied:

- EN 61010-1: 2001  
Safety requirements for electrical equipment for measurement, control and laboratory use -  
Part 1: General requirements
- EN 61010-2-101: 2002  
Safety requirements for electrical equipment for measurement, control and laboratory use -  
Part 2-101: Particular requirements for in vitro diagnostic (IVD) medical equipment
- EN 61010-2-010: 2003  
Safety requirements for electrical equipment for measurement, control and laboratory use -  
Part 2: Particular requirements for laboratory equipment for the heating of materials.
- EN 61326-1: 1997 + A1: 1998 + A2: 2001 + A3: 2003  
Electrical equipment for measurement, control and laboratory use -  
EMC requirements -  
Part 1: General requirements
- EN 61000-3-2: 2000  
Electromagnetic compatibility (EMC)  
Part 3-2: Limits - Limits for harmonic current emissions
- EN 61000-3-3: 1995 + A1: 2001  
Electromagnetic compatibility (EMC)  
Part 3: Limits -  
Section 3: Limitation of voltage fluctuations and flicker in low-voltage  
supply systems for equipment with rated current  $\leq 16$  A

In addition, the following in-house standards were applied:

- DIN EN ISO 9001: 2000.

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